



(19)

Europäisches Patentamt
European Patent Office
Office européen des brevets



(11)

EP 1 052 654 A1

(12)

EUROPEAN PATENT APPLICATION

(43) Date of publication:

15.11.2000 Bulletin 2000/46

(51) Int. Cl.⁷: H01B 3/44, C08K 7/06,
C08K 3/04, H01B 7/18

(21) Application number: 00303933.6

(22) Date of filing: 10.05.2000

(84) Designated Contracting States:

AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU
MC NL PT SE

Designated Extension States:

AL LT LV MK RO SI

(30) Priority: 13.05.1999 US 311480

(71) Applicant:

UNION CARBIDE CHEMICALS & PLASTICS
TECHNOLOGY CORPORATION
Danbury, Connecticut 06817-0001 (US)

(72) Inventors:

- Lee, Wei-kuo
Bridgewater, New Jersey 07080 (US)
- Avgousti, Marios
Plainsboro, New Jersey 08536 (US)

(74) Representative:

Hayes, Adrian Chetwynd et al
Boult Wade Tenant,
Verulam Gardens
70 Gray's Inn Road
London WC1X 8BT (GB)

(54) Cable semiconducting shield

(57) A cable comprising one or more electrical conductors or communications media or a core of two or more electrical conductors or communications media, each electrical conductor, communications medium, or core being surrounded by a layer comprising:

- (a) polyethylene; polypropylene; or mixtures thereof;
- (b) carbon nanotubes;
- (c) optionally, a conductive carbon black other than carbon nanotubes; and
- (d) optionally, a copolymer of acrylonitrile and butadiene wherein the acrylonitrile is present in an amount of about 30 to about 60 percent by weight based on the weight of the copolymer or a silicone rubber.

Description**Technical Field**

5 [0001] This invention relates mainly to a power cable having a semiconducting shield.

Background Information

10 [0002] A typical electric power cable generally comprises one or more conductors in a cable core that is surrounded by several layers of polymeric materials including a first semiconducting shield layer (conductor or strand shield), an insulating layer, a second semiconducting shield layer (insulation shield), a metallic tape or wire shield, and a protective jacket. The outer semiconducting shield can be either bonded to the insulation or strippable, with most applications using strippable shields. Additional layers within this construction such as moisture impervious materials are often incorporated.

15 [0003] Polymeric semiconducting shields have been utilized in multilayered power cable construction for many decades. Generally, they are used to fabricate solid dielectric power cables rated for voltages greater than 1 kilo Volt (kV). These shields are used to provide layers of intermediate conductivity between the high potential conductor and the primary insulation, and between the primary insulation and the round or neutral potential. The volume resistivity of these semiconducting materials is typically in the range of 10^1 to 10^8 ohm-cm when measured on a completed power cable 20 construction using the methods described in ICEA S-66-524, section 6.12, or IEC 60502-2 (1997), Annex C. Typical strippable shield compositions contain a polyolefin, such as ethylene/vinyl acetate copolymer with a high vinyl acetate content, conductive carbon black, an organic peroxide crosslinking agent, and other conventional additives such as a nitrile rubber, which functions as a strip force reduction aid, processing aids, and antioxidants. These compositions are usually prepared in granular or pellet form. Polyolefin formulations such as these are disclosed in United States patent 25 4,286,023 and European Patent Application 420 271. The shield composition is, typically, introduced into an extruder where it is co-extruded around an electrical conductor at a temperature lower than the decomposition temperature of the organic peroxide to form a cable. The cable is then exposed to higher temperatures at which the organic peroxide decomposes to provide free radicals, which crosslink the polymer.

30 [0004] In order to provide a semiconducting shield it is necessary to incorporate conductive particles into the composition. These conductive particles have been generally provided by particulate carbon black. Useful carbon blacks can have a surface area of about 50 to about 1000 square meters per gram. The surface area is determined under ASTM D 4820-93a (Multipoint B.E.T. Nitrogen Adsorption). The carbon blacks have been used in the semiconducting shield composition in an amount of about 20 to about 60 percent by weight based on the weight of the composition, and are preferably used in an amount of about 25 to about 45 percent by weight. Both standard conductivity and high conductivity carbon blacks are used with standard conductivity blacks being preferred. Examples of conductive carbon 35 blacks are the grades described by ASTM N550, N472, N351, N110, Ketjen blacks, and acetylene blacks.

35 [0005] Industry is constantly attempting to select carbon blacks, which, at a moderate cost, will improve cable strength and provide more efficient conductivity.

40 Disclosure of the Invention

[0006] An object of this invention, therefore, is to provide a cable having an improved semiconducting shield in terms of structure and conductivity. Other objects and advantages will become apparent hereinafter.

45 [0007] According to the invention, such a cable has been discovered. The cable comprises one or more electrical conductors or communications media or a core of two or more electrical conductors or communications media, each electrical conductor, communications medium, or core being surrounded by a layer comprising:

- (a) polyethylene; polypropylene; or mixtures thereof;
- (b) carbon nanotubes;
- 50 (c) optionally, a conductive carbon black other than carbon nanotubes; and
- (d) optionally, a copolymer of acrylonitrile and butadiene wherein the acrylonitrile is present in an amount of about 30 to about 60 percent by weight based on the weight of the copolymer or a silicone rubber.

55 Description of the Preferred Embodiment(s)

[0008] Polyethylene, as that term is used herein, is a homopolymer of ethylene or a copolymer of ethylene and a minor proportion of one or more alpha-olefins having 3 to 12 carbon atoms, and preferably 4 to 8 carbon atoms, and,

optionally, a diene, or a mixture or blend of such homopolymers and copolymers. The mixture can be a mechanical blend or an in situ blend of two or more polymers. Examples of the alpha-olefins are propylene, 1-butene, 1-hexene, 4-methyl-1-pentene, and 1-octene. The polyethylene can also be a copolymer of ethylene and an unsaturated ester such as a vinyl ester, e.g., vinyl acetate or an acrylic or methacrylic acid ester, which are preferably used in semiconducting shields.

[0009] The polyethylene can be homogeneous or heterogeneous. The homogeneous polyethylenes usually have a polydispersity (Mw/Mn) in the range of about 1.5 to about 3.5 and an essentially uniform comonomer distribution, and are characterized by single and relatively low DSC melting points. The heterogeneous polyethylenes, on the other hand, have a polydispersity (Mw/Mn) greater than 3.5 and do not have a uniform comonomer distribution. Mw is defined as weight average molecular weight and Mn is defined as number average molecular weight. The polyethylenes can have a density in the range of 0.860 to 0.950 gram per cubic centimeter, and preferably have a density in the range of 0.870 to about 0.930 gram per cubic centimeter. They also can have a melt index in the range of about 0.1 to about 50 grams per 10 minutes.

[0010] The polyethylenes can be produced by low or high pressure processes. They are preferably produced in the gas phase, but they can also be produced in the liquid phase in solutions or slurries by conventional techniques. Low pressure processes are typically run at pressures below 1000 psi (pounds per square inch) whereas high pressure processes are typically run at pressures above 15,000 psi.

[0011] Typical catalyst systems, which can be used to prepare these polyethylenes, are magnesium/titanium based catalyst systems, which can be exemplified by the catalyst system described in United States patent 4,302,565 (heterogeneous polyethylenes); vanadium based catalyst systems such as those described in United States patents 4,508,842 (heterogeneous polyethylenes) and 5,332,793; 5,342,907; and 5,410,003 (homogeneous polyethylenes); a chromium based catalyst system such as that described in United States patent 4,101,445; a metallocene catalyst system such as those described in United States patents 4,937,299, 5,272,236, 5,278,272, and 5,317,036 (homogeneous polyethylenes); or other transition metal catalyst systems. Many of these catalyst systems are often referred to as Ziegler-Natta catalyst systems or Phillips catalyst systems. Catalyst systems, which use chromium or molybdenum oxides on silica-alumina supports, can be included here. Typical processes for preparing the polyethylenes are also described in the aforementioned patents. Typical in situ polyethylene blends and processes and catalyst systems for providing same are described in United States Patents 5,371,145 and 5,405,901. The various polyethylenes can include low density homopolymers of ethylene made by high pressure processes (HP-LDPEs), linear low density polyethylenes (LLD-PEs), very low density polyethylenes (VLDPEs), medium density polyethylenes (MDPEs), high density polyethylene (HDPE) having a density greater than 0.940 gram per cubic centimeter and metallocene copolymers with densities less than 0.900 gram per cubic centimeter. The latter five polyethylenes are generally made by low pressure processes. A conventional high pressure process is described in Introduction to Polymer Chemistry, Stille, Wiley and Sons, New York, 1962, pages 149 to 151. The high pressure processes are typically free radical initiated polymerizations conducted in a tubular reactor or a stirred autoclave. In the stirred autoclave, the pressure is in the range of about 10,000 to 30,000 psi and the temperature is in the range of about 175 to about 250 degrees C, and in the tubular reactor, the pressure is in the range of about 25,000 to about 45,000 psi and the temperature is in the range of about 200 to about 350 degrees C.

[0012] HP-LDPE and the copolymers of ethylene and unsaturated esters are generally made by these high pressure processes. Blends of high pressure polyethylene and metallocene resins can also be used, the former component because of its excellent processability and the latter because of its flexibility.

[0013] As noted, the copolymers comprised of ethylene and unsaturated esters can be prepared by the conventional high pressure techniques described above and are preferred for semiconducting shields. The unsaturated esters can be alkyl acrylates, alkyl methacrylates, and vinyl carboxylates. The alkyl group can have 1 to 8 carbon atoms and preferably has 1 to 4 carbon atoms. The carboxylate group can have 2 to 8 carbon atoms and preferably has 2 to 5 carbon atoms. In semiconducting shields, the portion of the copolymer attributed to the ester comonomer can be in the range of about 20 to about 55 percent by weight based on the weight of the copolymer, and is preferably in the range of about 35 to about 55 percent by weight. The ester can have about 4 to about 20 carbon atoms, and preferably has about 4 to about 7 carbon atoms. Examples of vinyl esters (or carboxylates) are vinyl acetate, vinyl butyrate, vinyl pivalate, vinyl neanonanoate, vinyl neodecanoate, and vinyl 2-ethylhexanoate. Vinyl acetate is preferred. Examples of acrylic and methacrylic acid esters are lauryl methacrylate; myristyl methacrylate; palmityl methacrylate; stearyl methacrylate; 3-methacryloxy-propyltrimethoxysilane; 3-methacryloxypropyltrimethoxysilane; cyclohexyl methacrylate; n-hexylmethacrylate; isodecyl methacrylate; 2-methoxyethyl methacrylate; tetrahydrofurfuryl methacrylate; octyl methacrylate; 2-phenoxyethyl methacrylate; isobornyl methacrylate; isoctylmethacrylate; octyl methacrylate; isoctyl methacrylate; oleyl methacrylate; ethyl acrylate; methyl acrylate; t-butyl acrylate; n-butyl acrylate; and 2-ethylhexyl acrylate. Methyl acrylate, ethyl acrylate, and n- or t-butyl acrylate are preferred. The alkyl group can be substituted with an oxyalkyltrialkoxysilane, for example. The copolymers can have a density in the range of 0.900 to 0.990 gram per cubic centimeter, and preferably have a density in the range of 0.920 to 0.970 gram per cubic centimeter. The copolymers can

also have a melt index in the range of about 0.1 to about 100 grams per 10 minutes, and preferably have a melt index in the range of about 1 to about 50 grams per 10 minutes. One process for the preparation of a copolymer of ethylene and an unsaturated ester is described in United States Patent 3,334,081.

[0014] The VLDPE can be a copolymer of ethylene and one or more alpha-olefins having 3 to 12 carbon atoms and

5 preferably 3 to 8 carbon atoms. The density of the VLDPE can be in the range of 0.870 to 0.915 gram per cubic centimeter. It can be produced, for example, in the presence of (i) a catalyst containing chromium and titanium, (ii) a catalyst

containing magnesium, titanium, a halogen, and an electron donor; or (iii) a catalyst containing vanadium, an electron

donor, an alkyl aluminum halide modifier, and a halocarbon promoter. Catalysts and processes for making the VLDPE

10 are described, respectively, in United States patents 4,101,445; 4,302,565; and 4,508,842. The melt index of the

about 50 grams per 10 minutes. The portion of the VLDPE attributed to the comonomer(s), other than ethylene, can be

15 in the range of about 1 to about 49 percent by weight based on the weight of the copolymer and is preferably in the

range of about 15 to about 40 percent by weight. A third comonomer can be included, e.g., another alpha-olefin or a

15 diene such as ethylidene norbornene, butadiene, 1,4-hexadiene, or a dicyclopentadiene. Ethylene/propylene copoly-

mers and ethylene/propylene/diene terpolymers are generally referred to as EPRs and the terpolymer is generally

20 referred to as an EPDM. The third comonomer can be present in an amount of about 1 to 15 percent by weight based

on the weight of the copolymer and is preferably present in an amount of about 1 to about 10 percent by weight. It is

25 preferred that the copolymer contain two or three comonomers inclusive of ethylene.

[0015] The LLDPE can include the VLDPE and MDPE, which are also linear, but, generally, has a density in the

20 range of 0.916 to 0.925 gram per cubic centimeter. It can be a copolymer of ethylene and one or more alpha-olefins

having 3 to 12 carbon atoms, and preferably 3 to 8 carbon atoms. The melt index can be in the range of about 0.1 to

about 100 grams per 10 minutes, and is preferably in the range of about 1 to about 50 grams per 10 minutes. The alpha-

olefins can be the same as those mentioned above, and the catalysts and processes are also the same subject to var-

25 iations necessary to obtain the desired densities and melt indices.

[0016] As noted, included in the definition of polyethylene are homopolymers of ethylene made by a conventional

high pressure process. The homopolymer preferably has a density in the range of 0.910 to 0.930 gram per cubic cen-

timeter. The homopolymer can also have a melt index in the range of about 1 to about 5 grams per 10 minutes, and

25 preferably has a melt index in the range of about 0.75 to about 3 grams per 10 minutes.

[0017] Melt index is determined in accordance with ASTM D-1238, Condition E, measured at 190 degrees C.

[0018] The polyethylenes can be made moisture curable by making the resin hydrolyzable, which is accomplished

by adding hydrolyzable groups such as -Si(OR)₃ wherein R is a hydrocarbyl radical to the resin structure through copo-

lymerization or grafting. Suitable crosslinking agents are organic peroxides such as dicumyl peroxide; 2,5-dimethyl-2,5-

30 preferred. Hydrolyzable groups can be added, for example, by copolymerizing ethylene with an ethylenically unsatu-

rated compound having one or more -Si(OR)₃ groups such as vinyltrimethoxysilane, vinyltriethoxysilane, and gamma-

35 methacryloxypropyltrimethoxysilane or grafting these silane compounds to the resin in the presence of the aforemen-

tioned organic peroxides. The hydrolyzable resins are then crosslinked by moisture in the presence of a silanol conden-

30 sation catalyst such as dibutyltin dilaurate, dioctyltin maleate, dibutyltin diacetate, stannous acetate, lead naphthenate,

40 and zinc caprylate. Dibutyltin dilaurate is preferred. Examples of hydrolyzable copolymers and hydrolyzable grafted

copolymers are ethylene/vinyltrimethoxy silane copolymer, ethylene/gamma-methacryloxypropyltrimethoxy silane

density ethylene/1-butene copolymer, and vinyltrimethoxy silane grafted linear low density polyethylene.

[0019] In applications where moisture cured insulation is used, it is desirable to provide a moisture cured strippable

45 semiconducting shield to protect the insulation. The shield composition would then be prepared in the same manner as

[0020] With respect to polypropylene: homopolymers and copolymers of propylene and one or more other alpha-

olefins wherein the portion of the copolymer based on propylene is at least about 60 percent by weight based on the

50 weight of the copolymer can be used by themselves or in admixture with polyethylene. The polypropylene can be pre-

pared by conventional processes such as the process described in United States patent 4,414,132. The alpha-olefins

in the copolymer are preferably those having 2 or 4 to 12 carbon atoms.

[0021] Component (b) consists essentially of carbon nanotubes. Carbon nanotubes are made of carbon and are

55 high strength sub-micron sized fibril particles having a graphitic morphological structure and configuration (an entan-

gled three dimensional network). They have been referred to as carbon fibrils and graphite fibrils, and can be prepared

as described in United States patent 5,707,916.

[0022] A typical carbon nanotube can be described as a tube made up of eight layers of rolled-up graphite sheets

55 having a hollow core 0.005 micron in diameter and an outer diameter of 0.01 micron (100 Angstroms). The length of the

tube is 1 to 10 microns. Each of the graphite layers is made up of carbon atoms. The tube is not straight, but can be

acicular or serpentine. They usually appear as a plurality of intertwined or entangled fibers; are of high strength; and

have a high level of electrical conductivity. Graphite fibrils are vapor grown graphitic carbon nanotubes. They are produced as agglomerates somewhat like steel wool pads although, of course, on a much smaller scale. The carbon nanotubes have a black color and their composition is essentially pure carbon with a trace of residual metal oxide catalyst, which means that they can be considered clean with a low concentration of metal ions. Because of their porous structure, they have a very low bulk density, i.e., about 0.10 gram per cubic centimeter or 6.24 pounds per cubic foot; the BET surface area is about 250 square meters per gram; and the DBP Absorption is 450 cubic centimeter per 100 grams.

[0023] When the carbon nanotubes are essentially the only carbon in the semiconducting layer composition, they can be used in amounts of about 1 to about 35 parts by weight per 100 parts by weight of component (a), and are preferably used in amounts of 2 to about 20 parts by weight. When they are used together with another conductive carbon black, the weight ratio of carbon nanotubes to conductive carbon black can be about 0.1:1 to about 10:1, and the total of carbon nanotubes and other conductive carbon black can be in the range of about 5 to about 80 parts by weight per 100 parts by weight of component (a).

[0024] Component (c) is optional, and can be a conventional conductive carbon black commonly used in semiconducting shields. These carbon blacks are described above. Subject to the above weight ratios and parts by weight, the conductive carbon black other than carbon nanotubes can be used in amounts of about 13 to about 100 parts by weight per 100 parts by weight of component (a).

[0025] Component (d) is also optional. It can be a copolymer of acrylonitrile and butadiene wherein the acrylonitrile is present in an amount of about 30 to about 60 percent by weight based on the weight of the copolymer, and is preferably present in an amount of about 40 to about 50 percent by weight. This copolymer is also known as a nitrile rubber or an acrylonitrile/butadiene copolymer rubber. The density can be, for example, 0.98 gram per cubic centimeter and the Mooney Viscosity can be (ML 1+4) 50. Component (d) can also be a silicone rubber. The components are commonly used in semiconducting shields.

[0026] For each 100 parts by weight of component (a), i.e., polyethylene, polypropylene, or mixtures thereof, the other components can be present in about the following values (in parts by weight):

component		broad range	preferred range
(b)	carbon nanotubes	1 to 35	2 to 20
(c)	conductive carbon black (optional)	13 to 100	15 to 80
(d)	nitrile rubber* or silicone rubber (optional)	10 to 60	15 to 45
	weight ratio of (b) to (c)	1 to 10	3 to 8
		0.1:1 to 10:1	0.2:1 to 3:1

* nitrile rubber is a copolymer of acrylonitrile and butadiene.

[0027] Component (a) can be crosslinked. This is accomplished in a conventional manner with an organic peroxide or irradiation, the former being preferred. The amount of organic peroxide used can be in the range of about 0.15 to about 0.8 part by weight of organic peroxide for each 100 parts by weight of component (a), and is preferably in the range of about 0.3 to about 0.6 part by weight. Organic peroxide crosslinking temperatures can be in the range of about 130 to about 250 degrees C and are preferably in the range of about 140 to about 210 degrees C.

[0028] Examples of organic peroxides useful in crosslinking are dicumyl peroxide; t-butyl cumyl peroxide; lauroyl peroxide; benzoyl peroxide; tertiary butyl perbenzoate; di(tertiary-butyl) peroxide; cumene hydroperoxide; 2,5-dimethyl-2,5-di(t-butyl-peroxy)hexyne-3; 2,5-dimethyl-2,5-di(t-butyl-peroxy)hexane; tertiary butyl hydroperoxide; isopropyl percarbonate; and alpha,alpha'-bis(tertiary-butylperoxy)diisopropylbenzene.

[0029] Another form of crosslinking is by irradiation, typically by electron beam. The composition in pellet form is subjected to an electron beam at a given dose rate or exposed to a Gamma source of specific strength for a given period of time to impart a specific dose rate of radiation.

[0030] Conventional additives, which can be introduced into the composition, are exemplified by antioxidants, coupling agents, ultraviolet absorbers or stabilizers, antistatic agents, pigments, dyes, nucleating agents, reinforcing fillers or polymer additives, slip agents, plasticizers, processing aids, lubricants, viscosity control agents, tackifiers, anti-blocking agents, surfactants, extender oils, metal deactivators, voltage stabilizers, flame retardant fillers and additives, crosslinking agents, boosters, and catalysts, and smoke suppressants. Additives and fillers can be added in amounts ranging from less than about 0.1 to more than about 50 percent by weight based on the weight of the composition.

[0031] Examples of antioxidants are: hindered phenols such as tetrakis [methylene(3,5-di-tert-butyl-4-hydroxyhydrocinnamate)]methane, bis [(beta-(3,5-di-tert-butyl-4-hydroxybenzyl)-methylcarboxyethyl)]sulphide, 4,4'-thiobis(2-methyl-6-tert-butylphenol), 4,4'-thiobis(2-tert-butyl-5-methylphenol), 2,2'-thiobis(4-methyl-6-tert-butylphenol), and thiophenyl)phosphite and di-tert-butylphenyl-phosphonite; thio compounds such as dilaurylthiodipropionate, dimyristylthiodipropionate, and distearylthiodipropionate; various sioxanes; and various amines such as polymerized 2,2,4-trimethyl-1,2-dihydroquinoline, 4,4'-bis(alpha,alpha-demethylbenzyl)diphenylamine, and alkylated diphenylamines. Antioxidants can be used in amounts of about 0.1 to about 5 percent by weight based on the weight of the composition.

[0032] Compounding can be effected in a conventional melt/mixer or in a conventional extruder, and these terms are used in this specification interchangeably. Generally, the conductive shield composition is prepared in a melt/mixer and then pelletized using a pelletizer attachment or an extruder adapted for pelletizing. Both the melt/mixer, as the name implies, and the extruder, in effect, have melting and mixing zones although the various sections of each are known to those skilled in the art by different names. The semiconducting shield composition of the invention can be prepared in various types of melt/mixers and extruders such as a Brabender™ mixer, Banbury™ mixer, a roll mill, a Buss™ co-kneader, a biaxial screw kneading extruder, and single or twin screw extruders. A description of a conventional extruder can be found in United States patent 4,857,600. In addition to melt/mixing, the extruder can coat a wire or a core of wires. An example of co-extrusion and an extruder therefor can be found in United States patent 5,575,965. A typical extruder has a hopper at its upstream end and a die at its downstream end. The hopper feeds into a barrel, which contains a screw. At the downstream end, between the end of the screw and the die, is a screen pack and a breaker plate. The screw portion of the extruder is considered to be divided up into three sections, the feed section, the compression section, and the metering section, and two zones, the back heat zone and the front heat zone, the sections and zones running from upstream to downstream. In the alternative, there can be multiple heating zones (more than two) along the axis running from upstream to downstream. If it has more than one barrel, the barrels are connected in series. The length to diameter ratio of each barrel is in the range of about 15:1 to about 30:1. In wire coating, where the material is crosslinked after extrusion, the die of the crosshead feeds directly into a heating zone, and this zone can be maintained at a temperature in the range of about 120°C to about 260°C, and preferably in the range of about 140°C to about 220°C.

[0033] The advantages of the invention are as follows: Where the carbon is essentially carbon nanotubes, interface roughness between the insulation and the semiconducting shield is eliminated and the cleanliness of the semiconducting shield is increased. Where a combination of carbon nanotubes and another conductive carbon black is used, there is a lower total cost; a cleaner composition; lower filler loadings; higher production rates; easier manufacturing and end use compounding; and better mechanical and electrical properties. The basis for the advantages of the combination is that 1 part by weight of carbon nanotubes provides conductivity roughly equal to about 7.5 parts by weight of conductive carbon black. Further, there can be a synergistic effect between the carbon nanotubes and the conductive carbon blacks with respect to electrical properties, particularly conductivity, which is found to change less with time than systems containing only conductive carbon blacks, and it appears that the carbon nanotube blends are more stable. There is also a benefit with respect to rheological properties in terms of lower shear viscosity, which can lower power needs in compounding; improve processibility; and lower extrusion temperatures resulting in better thermal stability. In the mixed system, volume resistivity is adequate at lower viscosities, and shows very small change with temperature, which is advantageous for lower dissipation factor. It is expected that the carbon nanotubes will disperse better than the Ketjen blacks.

[0034] Carbon nanotubes can also be used in the insulation layer in amounts of about 0.01 to about 1 part by weight per 100 parts by weight of component (a), i.e., polyethylene, polypropylene, or mixtures thereof, and is preferably used in the insulation layer in amounts of about 0.05 to about 0.3 part by weight. The advantages of using carbon nanotubes in the insulation layer are reduction or prevention of water treeing and increasing breakdown strength by dissipating electric energy or reducing electric stress.

[0035] The term "surrounded" as it applies to a substrate being surrounded by an insulating composition, jacketing material, semiconducting shield, or other cable layer is considered to include extruding around the substrate; coating the substrate; or wrapping around the substrate as is well known by those skilled in the art. The substrate can include, for example, a core including a conductor or a bundle of conductors, or various underlying cable layers as noted above.

[0036] All molecular weights mentioned in this specification are weight average molecular weights unless otherwise designated.

[0037] The patents mentioned in this specification are incorporated by reference herein.

[0038] The invention is illustrated by the following examples:

Examples 1 to 4

[0039] Products suitable for semiconductive shielding and of electrical power cables are prepared at various ratios of conductive carbon black and carbon nanotubes. The viscosity and electrical properties of these compositions are then measured.

[0040] Example 1 is a material which is known in the art to be useful for semiconductive shielding of electrical power cables. This material is made from 60 weight percent, with respect to the weight of the composition, of ethylene/ethyl acrylate which is a copolymer of ethylene and ethyl acrylate having an 18 percent by weight ethylene acrylate content and a melt index of 20 grams per 10 minutes. Example 1 also contains 38 weight percent of commercially available conductive carbon black, Denka Granules, which is an acetylene black having a surface area of 60 square meters per gram determined under ASTM D 4820-93a (Multipoint B.E.T. Nitrogen Adsorption). Polymerized 2,2,4-trimethyl-1,2-dihydroquinoline is an antioxidant added at 1 weight percent, and the processing additive, polyethylene glycol, is also added at 1 weight percent. The composition of Example 1 is shown in Table 1.

[0041] Example 2 is a commercially available mixture of carbon nanotubes with polyethylene. The composition contains 78 weight percent, with respect to the weight of the composition, of a linear low density polyethylene (LLDPE) having a density of 0.92 gram per cubic centimeter and a melt index of 20 grams per 10 minutes. Example 2 also contains 20 weight percent of carbon nanotubes. This composition also contains approximately 1 weight percent of antioxidant. The composition of Example 2 is shown in Table 1.

[0042] Examples 3 and 4 are prepared by blending the compositions of Examples 1 and 2 with a 30 millimeter laboratory scale twin screw mixing device. Example 3 is a blend of Examples 1 and 2 in a ratio of 75:25. Example 4 is a blend of Examples 1 and 2 in a ratio of 50:50. The compositions of Examples 3 and 4 are shown in Table 1.

[0043] The apparent shear viscosities of Examples 1 to 4 are measured with a piston driven capillary rheometer, Göttfert Rheograph™ model 2001. The capillary die is 1 millimeter in diameter, and 20 millimeters in length. The test temperature is 125 degrees C, which is similar to the temperature at which commercial peroxide crosslinkable semiconductive materials are extruded for shielding of electrical power cables. The piston velocity is varied to obtain an apparent shear rate from 90 to 900 sec⁻¹, also a representative range for commercial extrusion processes used for semiconductive shielding products. The pressure drop across the capillary die is measured with a single pressure transducer. The viscosity is calculated from die dimensions, the piston cross section, piston velocity, and the pressure drop across the die. The results of these measurements are compiled in Table 1.

30

TABLE 1

Example Number	1	2	3	4
Composition, weight percent				
Ethylene/Ethyl Acrylate	60.0	0.0	45.0	30.0
LLDPE	0.0	79.0	19.8	39.5
Carbon Black	38.0	0.0	28.5	19.0
Carbon Nanotubes	0.0	20.0	5.0	10.0
Antioxidant	1.0	1.0	1.0	1.0
Process aid	1.0	0.0	0.8	0.5
Total:	100.0	100.0	100.0	100.0
Viscosity, Pascal*seconds				
at 90 sec ⁻¹ shear rate	3470	4480	1770	2520
at 180 sec ⁻¹ shear rate	2360	2940	1170	1560
at 360 sec ⁻¹ shear rate	1570	1810	840	1000
at 900 sec ⁻¹ shear rate	858	875	506	557

The viscosity of Example 1 at 360 sec⁻¹ shear rate, 1570 Pascal seconds, is very typical for commercial products useful

EP 1 052 654 A1

as semiconductive shielding for power cables. This viscosity is dictated by the dual requirements of high molecular weight to maintain adequate mechanical properties, and sufficient carbon black to ensure adequately low volume resistivity. The viscosities of Example 3 and Example 4 are significantly lower than Example 1, which is advantageous for the manufacturing of insulated power cables.

5

Examples 5 to 8

10

[0044] Examples 5 to 8 are prepared by adding 1.1 weight percent, with respect to the weight of the composition, of dicumyl peroxide. The materials are then compression molded into slabs with 170 megapascals applied pressure, and then cured for 15 minutes at 175 degrees C. The molded slabs are cut into rectangular samples with dimensions approximately 3 mm thick, 25 mm wide, and 70 mm in length. Conductive silver paint, (DuPont™ grade 4817N) is used to apply electrodes on these samples across the width and thickness, separated by a distance of 50 millimeters. After curing, electrical leads are connected to the silver electrodes with clips. The resistance of the samples is measured with a two wire resistance meter. The volume resistivity is calculated from the measured resistance, the length between the electrodes (50 mm), and the cross sectional area of the sample (75 mm²). Resistance is measured at high temperatures by placing the sample inside a laboratory oven.

15

20

25

[0045] The volume resistivity of Examples 5 to 8 is measured at room temperature, and at 90 and 130 degrees C after exposure in the oven for 1 and 15 days. After 15 days, the samples are removed from the oven and allowed to cool for one day at room temperature, at which time the resistance is measured again. The results of these measurements on Examples 5 to 8 are shown in Table 2. The coefficient of variation (i.e., standard deviation divided by the mean result) for this test method is approximately 10 percent. Relative differences less than 20 percent are not significant.

TABLE 2

Example Number	5	6	7	8
Composition, weight percent				
Example 1	98.9			
Example 2		98.9		
Example 3			98.9	
Example 4				98.9
dicumyl peroxide	1.1	1.1	1.1	1.1
Total:	100.0	100.0	100.0	100.0
Volume Resistivity (ohm centimeters)				
at 23°C	13	4.0	67	14
at 90°C, 1 day exposure	50	4.2	150	19
at 90°C, 15 days exposure	47		150	20
at 23°C, after 15 days at 90°C	13		82	17
at 130°C, 1 day exposure	260	5.6	220	18
at 130°C, 15 days exposure	250		200	18
at 23°C, after 15 days at 130°C	28		110	17

55

[0046] The volume resistivity of the semiconductive shield on insulated power cable should be as low as possible. The dielectric loss factor of the cable, when used for AC electrical power transmission at voltages greater than 5 kV, is related to the volume resistivity of the shield layers. As the resistivity is increased, the dissipation factor also increases. It is desirable to minimize the dielectric loss of the power cable, and hence, it is desirable to minimize the volume resistivity.

[0047] Example 5 shows the volume resistivity for materials which are used commercially to shield power cables. The volume resistivity of the crosslinked semiconductive material increases with temperature, and remains stable after exposure to high temperatures for an extended period of time. After exposure to 130 degrees C and cooling down to room temperature, this material exhibits a significant increase in volume resistivity, which is undesirable.

5 [0048] Example 7 exhibits behavior similar to Example 5 except that there is less of a relative increase in volume resistivity with temperature, and less relative permanent increase in the room temperature volume resistivity after the temperature cycle. Even though the absolute value of volume resistivity for Example 7 is higher than Example 5, the stability of volume resistivity with temperature cycling and time is very desirable.

10 [0049] Example 8 exhibits superior volume resistivity characteristics relative to Example 5. This semiconductive composition exhibits very little temperature dependence on volume resistivity, and essentially no change in volume resistivity after temperature cycling.

[0050] The lower absolute volume resistivity and improved thermal stability of volume resistivity of Example 8 is highly desirable behavior, and is unexpected, especially when viewed in light of the lower viscosity measured on this composition in Example 4.

15

Claims

1. A cable comprising one or more electrical conductors or communications media or a core of two or more electrical conductors or communications media, each electrical conductor, communications medium, or core being surrounded by a layer comprising:

- (a) polyethylene; polypropylene; or mixtures thereof;
- (b) carbon nanotubes;
- (c) optionally, a conductive carbon black other than carbon nanotubes; and
- (d) optionally, a copolymer of acrylonitrile and butadiene wherein the acrylonitrile is present in an amount of 30 to 60 percent by weight based on the weight of the copolymer or a silicone rubber.

2. The cable defined in claim 1 wherein component (a) is a copolymer of ethylene and an unsaturated ester.

30 3. The cable defined in claim 2 wherein the copolymer is selected from the group consisting of vinyl esters, acrylic acid esters, and methacrylic acid esters wherein the ester is present in the copolymer in an amount of 20 to 55 percent by weight.

35 4. The cable defined in any one of the preceding claims wherein the layer is a semiconducting shield and component (b) is present in an amount of 13 to 100 parts by weight per 100 parts by weight of component (a).

5. The cable defined in any one of the preceding claims wherein the layer is a semiconducting shield and, for each 100 parts of component (a), component (b) is present in an amount of 1 to 35 parts by weight; component (c) is present in an amount of 13 to 100 parts by weight; and the weight ratio of component (b) to component (c) is 0.1:1 to 10:1.

40 6. The cable defined in any one of claims 1 to 3 wherein the layer is an insulation layer and component (b) is present in an amount of 0.01 to 1 part by weight per 100 parts by weight of component (a).

45 7. A cable comprising one or more electrical conductors or communications media or a core of two or more electrical conductors or communications media, each electrical conductor, communications medium, or core being surrounded by a semiconducting shield layer comprising:

- (a) a copolymer selected from the group consisting of vinyl esters, acrylic acid esters, and methacrylic acid esters wherein the ester is present in the copolymer in an amount of 20 to 55 percent by weight;
- (b) carbon nanotubes;
- (c) a conductive carbon black other than carbon nanotubes; and
- (d) optionally, a copolymer of acrylonitrile and butadiene wherein the acrylonitrile is present in an amount of 30 to 60 percent by weight based on the weight of the copolymer or a silicone rubber;

55

with the conditions that, for each 100 parts of component (a), component (b) is present in an amount of 2 to 20 parts by weight; component (c) is present in an amount of 15 to 80 parts by weight; and the weight ratio of component (b) to component (c) is 0.2:1 to 8:1.

8. A cable comprising one or more electrical conductors or communications media or a core of two or more electrical conductors or communications media, each electrical conductor, communications medium, or core being surrounded by a layer comprising:

5 (a) polyethylene; polypropylene; or mixtures thereof; and
(b) carbon nanotubes

with the condition that, for each 100 parts of component (a), component (b) is present in an amount of 0.05 to 0.3 part by weight.

10 9. A composition comprising:

(a) a copolymer selected from the group consisting of vinyl esters, acrylic acid esters, and methacrylic acid esters wherein the ester is present in the copolymer in an amount of 20 to 55 percent by weight;
15 (b) carbon nanotubes;
(c) a conductive carbon black other than carbon nanotubes; and
(d) optionally, a copolymer of acrylonitrile and butadiene wherein the acrylonitrile is present in an amount of 30 to 60 percent by weight based on the weight of the copolymer or a silicone rubber
with the conditions that, for each 100 parts of component (a), component (b) is present in an amount of 1 to 35 parts by weight; component (c) is present in an amount of 13 to 100 parts by weight; and the weight ratio of component (b) to component (c) is 0.1:1 to 10:1.

20 10. A composition comprising a polymer selected from the group consisting of polyethylene; polypropylene; and mixtures thereof and 0.01 to 1 part by weight of carbon nanotubes per 100 parts by weight of polymer.

25

30

35

40

45

50

55



EUROPEAN SEARCH REPORT

Application Number
EP 00 30 3933

DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int.Cl.)
A	WO 90 12842 A (JAPAN SYNTHETIC RUBBER CO LTD ;HYPERION CATALYSIS INT (US)) 1 November 1990 (1990-11-01) * the whole document *	1-10	H01B3/44 C08K7/06 C08K3/04 H01B7/18
A	WO 90 10296 A (JAPAN SYNTHETIC RUBBER CO LTD ;HYPERION CATALYSIS INT (US)) 7 September 1990 (1990-09-07) * the whole document *	1-10	
D,A	EP 0 420 271 A (UNION CARBIDE CHEM PLASTIC) 3 April 1991 (1991-04-03) * the whole document *	1-10	
D,A	US 4 286 023 A (ONGCHIN LUCIO) 25 August 1981 (1981-08-25) * the whole document *	1-10	
TECHNICAL FIELDS SEARCHED (Int.Cl.)			
H01B C08K			
The present search report has been drawn up for all claims			
Place of search	Date of completion of the search		Examiner
THE HAGUE	18 July 2000		Shade, M
CATEGORY OF CITED DOCUMENTS		T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document	
X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document			

ANNEX TO THE EUROPEAN SEARCH REPORT
ON EUROPEAN PATENT APPLICATION NO.

EP 00 30 3933

This annex lists the patent family members relating to the patent documents cited in the above-mentioned European search report. The members are as contained in the European Patent Office EDP file on. The European Patent Office is in no way liable for these particulars which are merely given for the purpose of information.

18-07-2000

Patent document cited in search report		Publication date	Patent family member(s)		Publication date
WO 9012842	A	01-11-1990	JP	2276839 A	13-11-1990
			JP	2863192 B	03-03-1999
			AU	5535290 A	16-11-1990
			US	5304326 A	19-04-1994
WO 9010296	A	07-09-1990	JP	2229267 A	12-09-1990
			JP	2232244 A	14-09-1990
			JP	2235945 A	18-09-1990
			AU	5336190 A	26-09-1990
EP 0420271	A	03-04-1991	AT	116073 T	15-01-1995
			CA	2026472 A,C	30-03-1991
			DE	69015302 D	02-02-1995
			DE	69015302 T	18-05-1995
			JP	3127403 A	30-05-1991
US 4286023	A	25-08-1981	US	4246142 A	20-01-1981
			CA	1100306 A	05-05-1981